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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) CA SUBSCRIBER PRICE	SINCE FILE ENTRY -1.60	TOTAL SESSION -1.60
=> file registry COST IN U.S. DOLLARS FULL ESTIMATED COST	SINCE FILE ENTRY 11.35	TOTAL SESSION 259.57
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) CA SUBSCRIBER PRICE	SINCE FILE ENTRY -1.60	TOTAL SESSION -1.60

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STRUCTURE FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3
DICTIONARY FILE UPDATES: 21 JAN 2008 HIGHEST RN 1000370-19-3

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TSCA INFORMATION NOW CURRENT THROUGH June 29, 2007

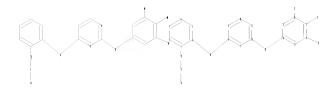
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http://www.cas.org/support/stngen/stndoc/properties.html

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19 20 21 22 23 24 25 26 rring nodes:
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 chain bonds:
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 chain bonds:
1-24 6-19 8-19 12-20 14-20 16-21 17-22 18-23 24-25 25-26 ring bonds:
1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 13-14 13-18 14-15 15-16 16-17 17-18 exact/norm bonds:
6-19 8-19 12-20 14-20 24-25 25-26 exact bonds:
1-24 16-21 17-22 18-23 normalized bonds:
1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 13-14 13-18 14-15 15-16 16-17 17-18
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Match level: 1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:CLASS 20:CLASS 21:CLASS 21:CLASS

L5 STRUCTURE UPLOADED

=> 15 sss ful

chain nodes :

L5 IS NOT A RECOGNIZED COMMAND

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=> s 15 sss ful

FULL SEARCH INITIATED 12:53:21 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 66 TO ITERATE

100.0% PROCESSED 66 ITERATIONS SEARCH TIME: 00.00.01

00.01

5 ANSWERS

L6 5 SEA SSS FUL L5

=> d 16 1-6

- ANSWER 1 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN 1.6
- 848468-37-1 REGISTRY RN
- ED Entered STN: 14 Apr 2005
- Benzenesulfonamide, N-propyl-2-[[2-[(3,4,5-trimethoxyphenyl)amino]-4
 - pyrimidinyl]amino]- (CA INDEX NAME)
- MF C22 H27 N5 O5 S
- SR CA
- LĊ STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- L6 ANSWER 2 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 848468-35-9 REGISTRY
- ED Entered STN: 14 Apr 2005
- CN Benzenesulfonamide, N,N-dimethyl-2-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-
- pyrimidinyl]amino]- (CA INDEX NAME)
- MF C21 H25 N5 O5 S
- CA SR
 - LĊ STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- ANSWER 3 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN L6
- RN 848468-34-8 REGISTRY
- Entered STN: 14 Apr 2005
- Benzenesulfonamide, N-(2-hydroxyethyl)-2-[[2-[(3,4,5-
- trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME) ME
- C21 H25 N5 06 S
- SR CA
- T.C STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L6 ANSWER 4 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 604802-45-1 REGISTRY
- ED Entered STN: 15 Oct 2003
- CN Benzenesulfonamide, N,2-dimethyl-6-[[2-[(3,4,5-trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)
- MF C21 H25 N5 O5 S
- SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L6 ANSWER 5 OF 5 REGISTRY COPYRIGHT 2008 ACS on STN
- RN 604802-41-7 REGISTRY
- ED Entered STN: 15 Oct 2003
- N Benzenesulfonamide, N-methyl-2-[[5-(trifluoromethyl)-2-[(3,4,5-
- trimethoxyphenyl)amino]-4-pyrimidinyl]amino]- (CA INDEX NAME)
- MF C21 H22 F3 N5 05 S SR CA
- LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file caplus COST IN U.S. DOLLARS SINCE FILE TOTAL SESSION ENTRY FULL ESTIMATED COST 188.82 448.39 DISCOUNT AMOUNTS (FOR OUALIFYING ACCOUNTS) SINCE FILE TOTAL SESSION ENTRY CA SUBSCRIBER PRICE 0.00 -1.60

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=> s 16 L7

2 L6

=> d 17 ibib abs 1-2

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:260035 CAPLUS

DOCUMENT NUMBER: 142:336377

TITLE: Preparation of 2,4-di(phenylamino)pyrimidines useful in the treatment of proliferative disorders

INVENTOR(S): Imbach, Patricia; Roesel, Johannes

PATENT ASSIGNEE(S): SOURCE: Novartis AG, Switz.; Novartis Pharma GmbH PCT Int. Appl., 39 pp. CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE:

GI

Patent English

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. WO 2005026130 A1 20050324 WO 2004-EP10466 20040917 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG AU 2004272288 A1 20050324 AU 2004-272288 20040917 CA 2538413 A1 20050324 CA 2004-2538413 20040917 EP 1663992 20060607 EP 2004-765358 20040917 A1 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK CN 1852900 20061025 CN 2004-80026942 Α 20040917 BR 2004014544 Α 20061107 BR 2004-14544 20040917 JP 2007505858 Т 20070315 JP 2006-526595 20040917 US 2007105839 A1 20070510 US 2006-571733 20060314 IN 2006CN00922 Α 20070615 IN 2006-CN922 20060316 MX 2006PA03054 MX 2006-PA3054 Α 20060531 US 2003-504374P PRIORITY APPLN. INFO.: 20030918 WO 2004-EP10466 W 20040917 OTHER SOURCE(S): CASREACT 142:336377; MARPAT 142:336377

AB The title compds. I [X = CRO, N; RO, R1-R4 = H, OH, alkyl, etc.; or R3 and R4 form together with the nitrogen and carbon atoms to which they are attached a 5-10 membered heterocyclic ring and comprising addnl. 1-3 heteroatoms selected from N, O and S; or R1-R3 = halo, haloalkyl, alkoxy, etc.; or R1 and R2 form aryl or 5-10 membered heteroaryl; R5, R6 = H, halo, CN, alkyl, etc.; R7-R9 = H, OH, alkyl, etc.], useful for preventing or treating proliferative disorders such as a tumor disease, by inhibiting ALK activity, were prepared E.g., a 2-step synthesis of 2-[2-(1H-indazol-6-ylamino)-pyrimidin-4-ylamino|benzenesulfonamide, starting from 2-aminobenzenesulfonamide and 2,4-dichloropyrimidine, was given. The compds. I were tested for inhibition of ALK tyrosine kinase in various cellular assays (data were given for representative compds. I).

Ι

L7 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:757684 CAPLUS

DOCUMENT NUMBER: 139:292258

TITLE: Pyrimidine derivatives

INVENTOR(S): Baenteli, Rolf; Zenke, Gerhard; Cooke, Nigel Graham; Duthaler, Rudolf; Thoma, Gebhard; Von Matt, Anette; Honda, Toshivuki; Matsuura, Naoko; Nonomura, Kazuhiko; Ohmori, Osamu; Umemura, Ichiro; Hinterding, Klaus;

Papageorgiou, Christos

PATENT ASSIGNEE(S): Novartis A.-G., Switz.; Novartis Pharma G.m.b.H. SOURCE:

PCT Int. Appl., 45 pp.

CODEN: PIXXD2 DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PA	PATENT NO.				KIND DATE			APPLICATION NO.					DATE					
WO	WO 2003078404			A1 20030925			WO 2003-EP2710				20030314							
											BG,							
											, EE							
											, KP							
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											A, US,							
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						FR,	GB,	GR,	HU,	IE	E, IT	LU,	MC,	NL,	PT,	RO,	SE,	
			SK,															
CA	CA 2479133			A1	20030925 CA 2003-247913				133	20030314								
AU	J 2003227070 A1				20030929 AU 2003-227070			70	20030314									
AU	2003227070			B2 20070201			EP 2003-744366											
EP																		
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BK	BR 2003008461				A 20050118					BR 2003-8461					20030314			
ON	JP 2005527529			20050915				ON 2003-376410					20030314					
NZ	BR 2003008461 JP 2005527529 CN 1697830 NZ 535109			A 20051110				NZ 2003-606101					20030314					
IN Z	IN 2004CN02025			7 20060326				TNI	2003	-0231	25	20030314						
	MX 2004PA09058 NO 2004004374																	
IIS	2006	1002	27		A1		2006	0511		IIS	2005	-5070	60		- 3	20050	613	
7.A	2004	0670	9		A		2006	0531		ZA	2004-	-6709	00		- 3	20060	330	
PRIORIT	Y APP	LN.	INFO	. :						GB	2004-	-6215			A :	20020		
											2003-					20030		
OTHER S	OURCE	(S):			MARI	PAT	139:	2922	58									

GI

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The pyrimidine derivs. (I) are claimed, wherein X = =CR or =N, R, R1, R2,
     R3, R4 independently is H, OH, C1-8alkyl, C2-8alkenyl, C3-8cycloalkyl,
     C3-8cycloalkyl-C1-8alkyl, hydroxyC1-8alkyl, C1-8alkoxyC1-8alkyl,
     hydroxyC1-8alkoxyC1-8alkyl, arylC1-8alkyl which optionally may be
     substituted on the ring by OH, C1-8alkoxy, carboxy, C1-8alkoxycarbonyl or
     R3 and R4 form together with N and C atoms to which they are attached to a
     5-10 membered heterocyclic ring containing 1, 2 or 3 heteroatoms of N, O or S;
     R1 and R2 form together with C atoms to which they are attached arvl of
     5-10 membered heteroarvl moiety containing 1-2 heteroatoms of N. O. S; R and
     R6 independently is H, halo, CN, C1-8alkyl, haloC1-8alkyl, C2-8alkenyl,
     C2-8alkynyl, C3-8cycloalkyl, C3-8cycloalkylC1-8alkyl, C5-10arylC1-8alkyl,;
     R7, R8 and R9 is independently H, OH, C1-8alkyl, C2-8alkenyl,
     haloC1-8alkyl, C1-8alkoxy, C3-8cycloalkyl, C3-8cycloalkylC1-8,
     arylC1-8alkyl. disorders where ZAP-70 and/or Syk inhibition plays a role
     or caused by a malfunction of signal cascades connected with FAK. I are
     useful in disorders where ZAP-70 and/or Syk inhibition plays a role or
     caused by a malfunction of signal cascades connected with FAK.
     Pharmaceutical compns. containing I are claimed.
REFERENCE COUNT:
                               THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
=> d his
     (FILE 'HOME' ENTERED AT 11:53:56 ON 22 JAN 2008)
     FILE 'REGISTRY' ENTERED AT 11:54:20 ON 22 JAN 2008
L1
               STRUCTURE UPLOADED
L2
              0 S L1 FAM FUL
L3
              5 S L1 SSS FUL
     FILE 'CAPLUS' ENTERED AT 11:55:10 ON 22 JAN 2008
              2 S L3
L4
     FILE 'REGISTRY' ENTERED AT 12:52:56 ON 22 JAN 2008
L5
                STRUCTURE UPLOADED
L6
              5 S L5 SSS FUL
     FILE 'CAPLUS' ENTERED AT 12:53:50 ON 22 JAN 2008
              2 S L6
=> logoff
ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF
LOGOFF? (Y)/N/HOLD:y
COST IN U.S. DOLLARS
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FULL ESTIMATED COST
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
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                                                       ENTRY
CA SUBSCRIBER PRICE
                                                        -1.60
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STN INTERNATIONAL LOGOFF AT 12:56:45 ON 22 JAN 2008

AB